





# The contribution of long-range transport and secondary organic aerosol to $PM_{2.5}$ in Pittsburgh

Juan C. Cabada-Amaya, Spyros N. Pandis, Allen L. Robinson, Ramachandran Subramanian, Wei Tang, Natalie J. Anderson, Timothy Raymond, Cliff I. Davidson

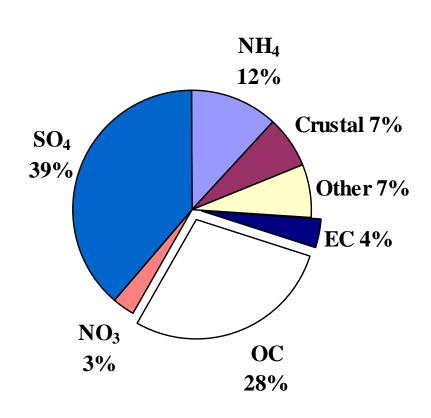
### Outline

- Motivation and Definitions.
- Sampling methods of Carbonaceous material at PAQS.
- Carbonaceous material composition for July 2001.
- Carbonaceous material composition from July 2001 to February 2002.





# PM<sub>2.5</sub> composition PAQS.



 $NH_4$ 10%  $SO_4$ Crustal 6% 30% **EC 4%**  $NO_3$ 10% OC 40%

July 2001,  $20 \,\mu g/m^3$ 

December 2001,  $10 \mu g/m^3$ 

### 18

### **OC/EC Sources**









Secondary Organic Aerosol Production in region



**Biogenic Aerosol** 



**Primary OC and EC Emissions in region** 



**Power Plants** 



**Automobiles** 



**Diesel Trucks** 



**Wood Burning** 



**Factories** 

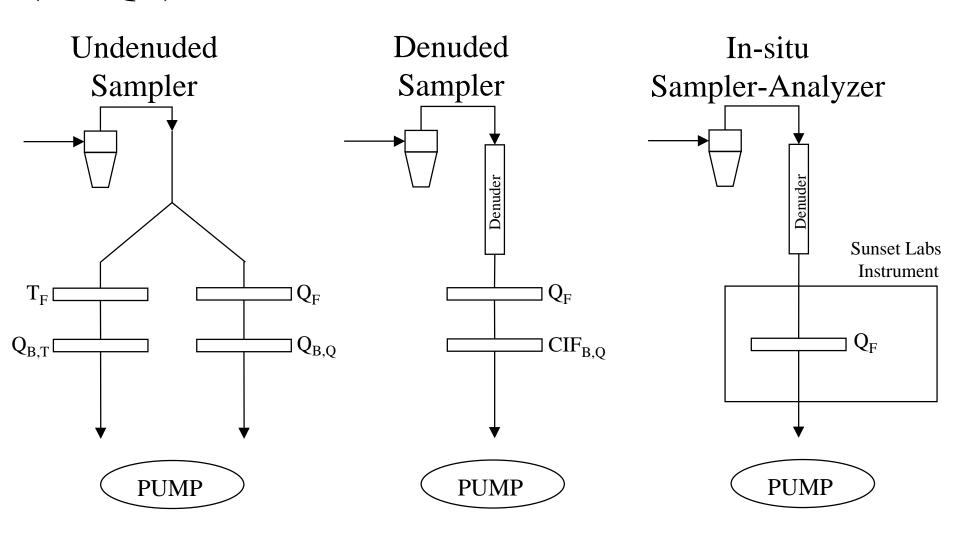


### PAQS Central sampling site



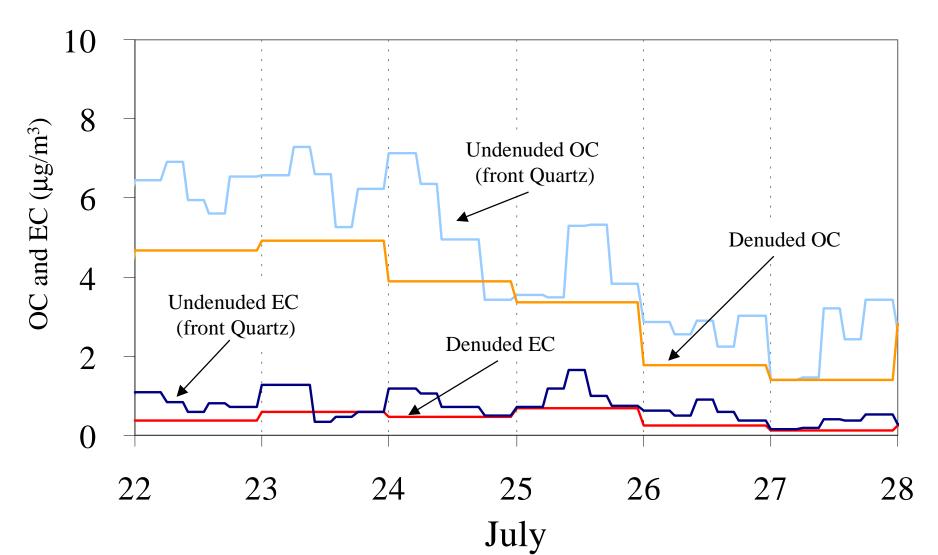


# OC and EC Sampler Configurations (PAQS)





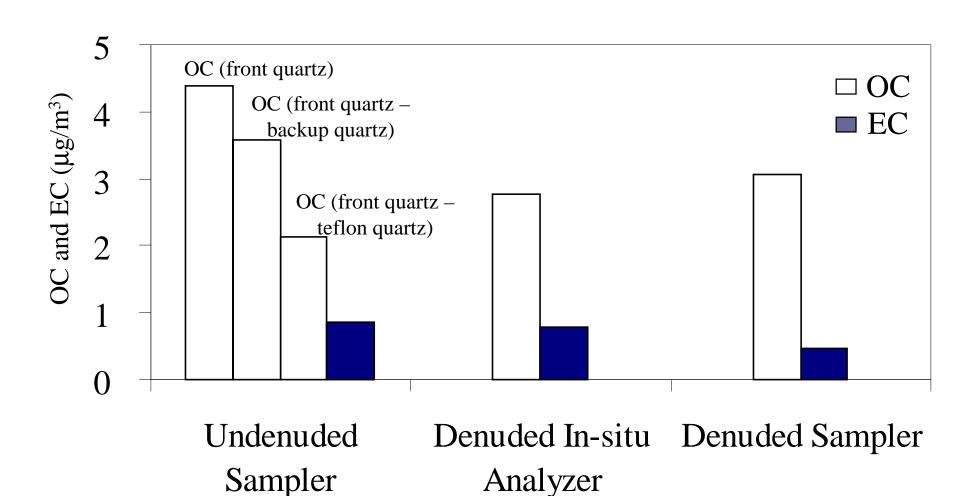
# OC and EC Measurements (PAQS, July 2001)







# Average OC and EC concentrations (PAQS, July 2001)



### Questions?

- Where is carbonaceous material coming from?
  - Regional or Local?
- What fraction of the measured organic carbon is primary and secondary?



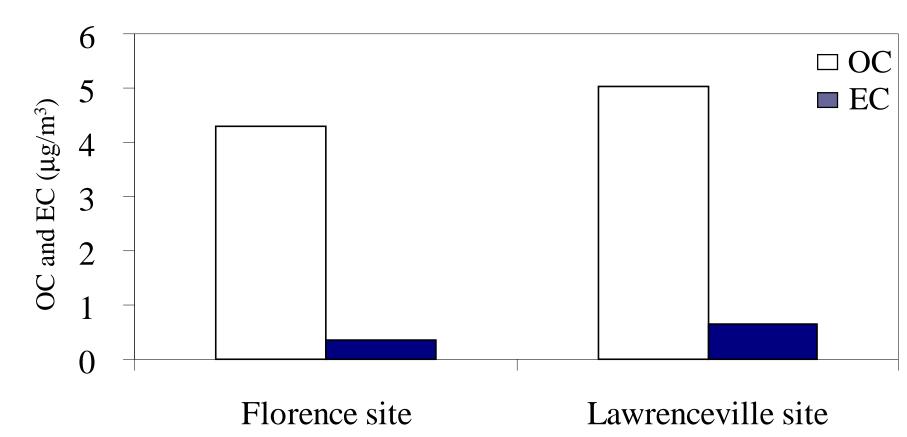
### Sampling Sites







# Regional Contribution of OC and EC to Pittsburgh



 Around 80% of carbonaceous material in Pittsburgh during July 2001 was due to long transport processes

### Primary vs. Secondary OC

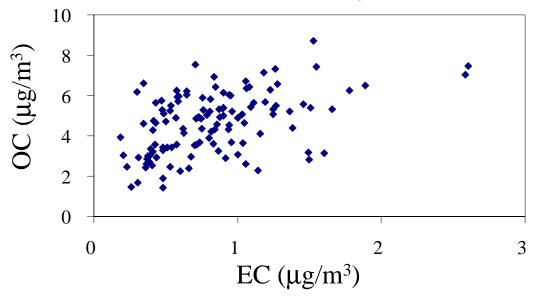
• EC can be used as tracer for primary OC.

$$OC|_{secondary} = OC|_{total} - EC|_{total} * OC/EC|_{emitted}$$

- Ambient Samples  $(OC|_{total})$  and  $EC|_{total}$
- Determine ratio of OC/EC primary emissions.
  (OC/EC | emitted)

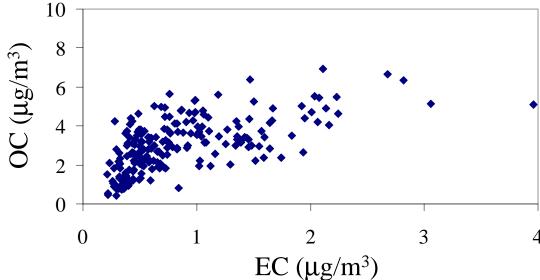


# OC and EC high resolution measurements, July 2001



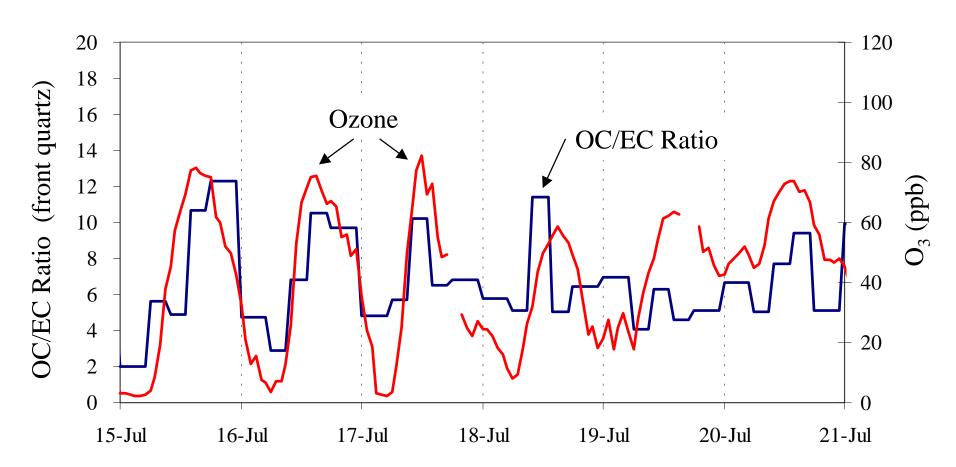
Undenuded Sampler (4-6 hrs samples)

Denuded In-situ Analyzer (2-4 hrs samples)

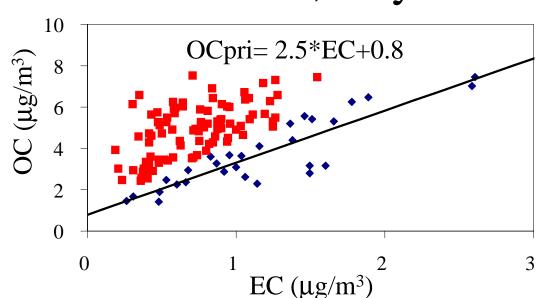




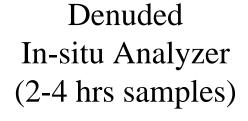
# Ozone as indicator of Photochemical activity

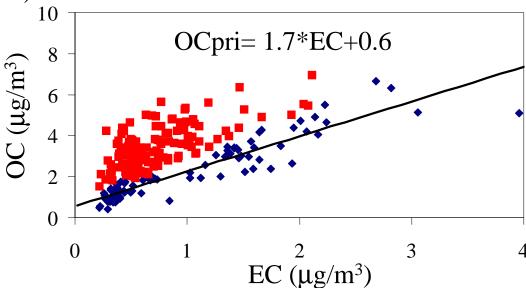


# OC/EC Ratio from measurements, July 2001



Undenuded Sampler (4-6 hrs samples)







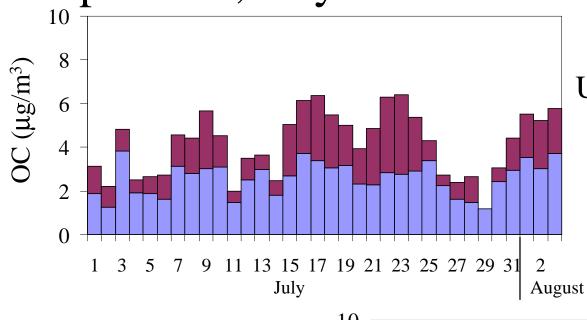
# Primary OC/EC Ratios from high resolution measurements, July 2001

Measurement (2-6 hrs)	OC/EC pri	Non- combustion primary OC
Undenuded (front quartz)	2.5	0.8
Undenuded (front quartz –teflon quartz correction)	1.0	0.4
Undenuded (front quartz - backup quartz correction)	2.5	0.1
Denuded In-situ analyzer	1.8	0.4

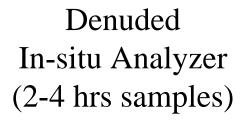
Based on NIOSH method measurements

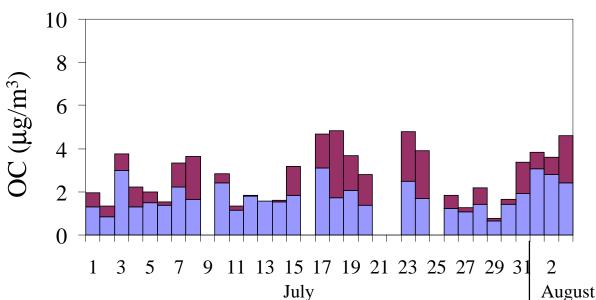
### CARNEGIE MELLON UNIVERSITY

# Daily Averaged OC Composition, July 2001



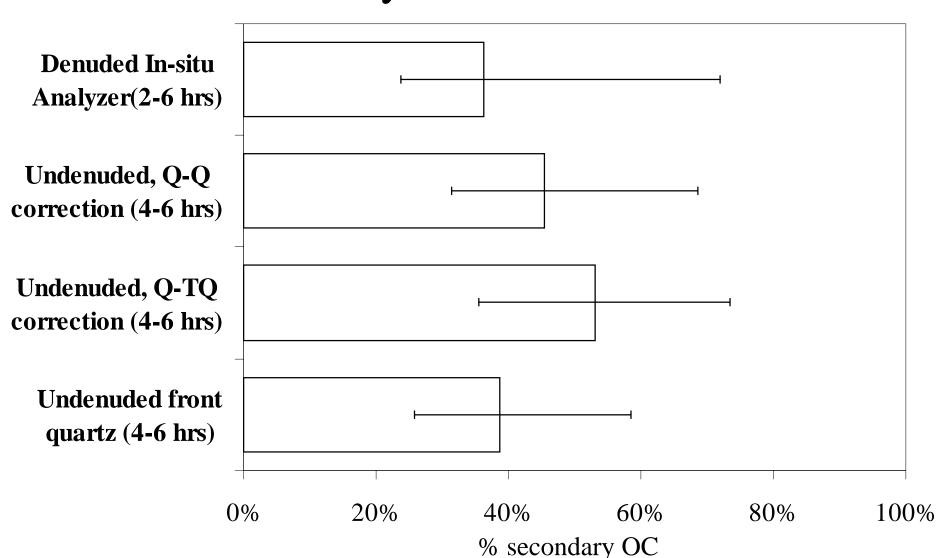
Undenuded Sampler (4-6 hrs samples)





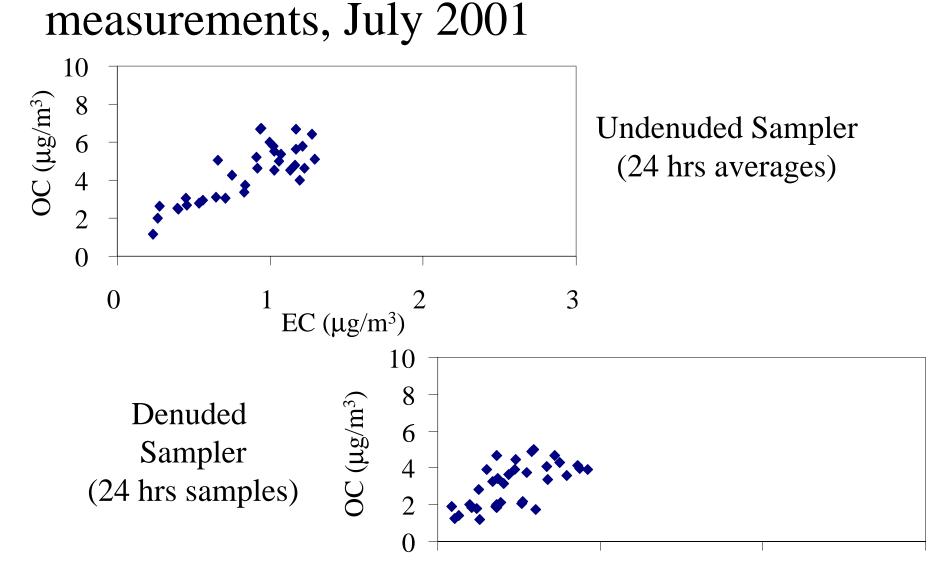


# OC Composition, high resolution measurements, July 2001

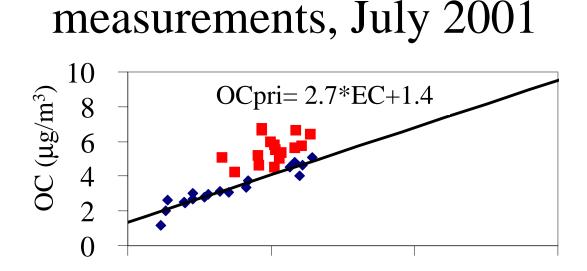


EC ( $\mu$ g/m<sup>3</sup>)

# OC and EC Daily

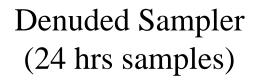


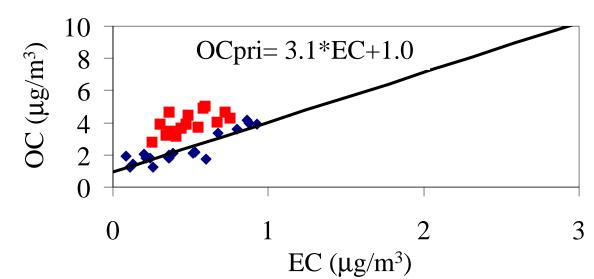
## OC/EC Ratio from



EC ( $\mu$ g/m<sup>3</sup>)

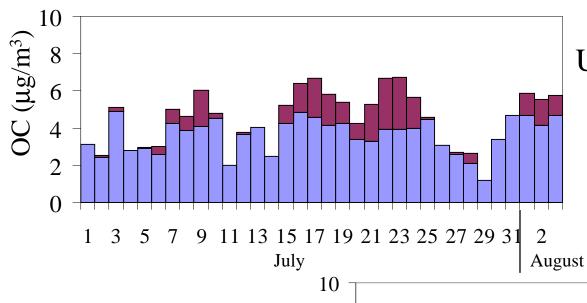
Undenuded Sampler (24 hrs averages)



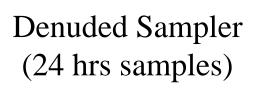


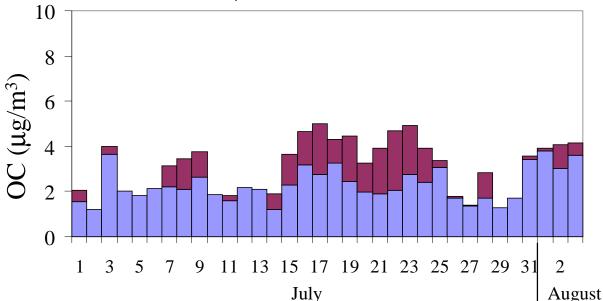


### Daily OC Composition, July 2001

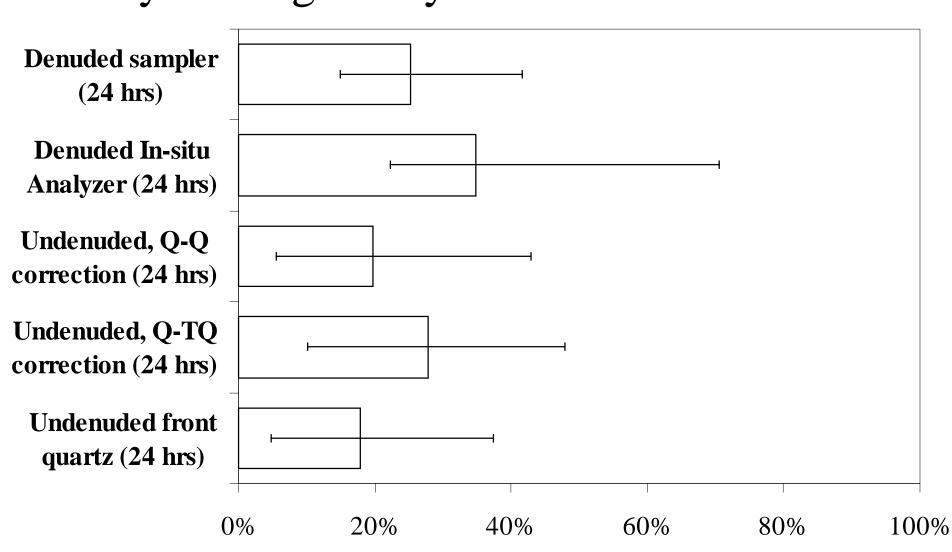


**Undenuded Sampler** (24 hrs averages)







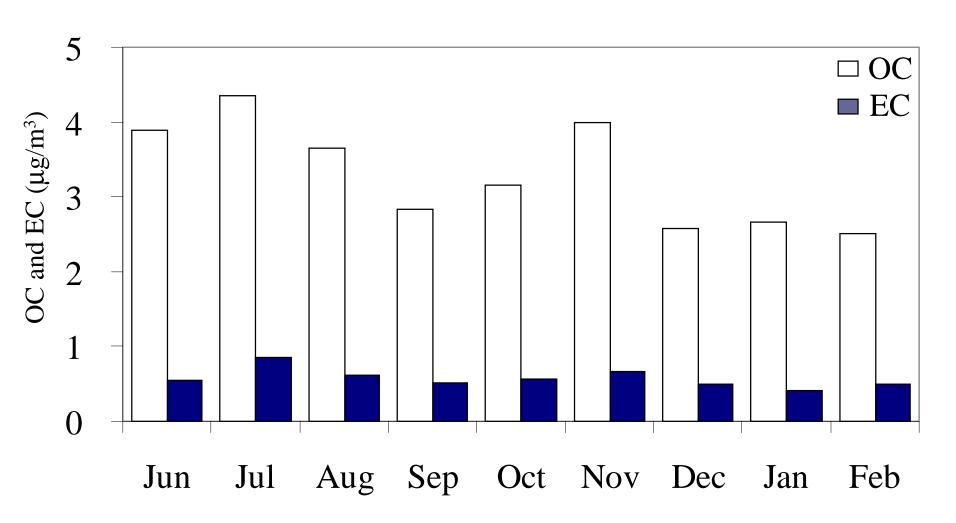


% secondary OC





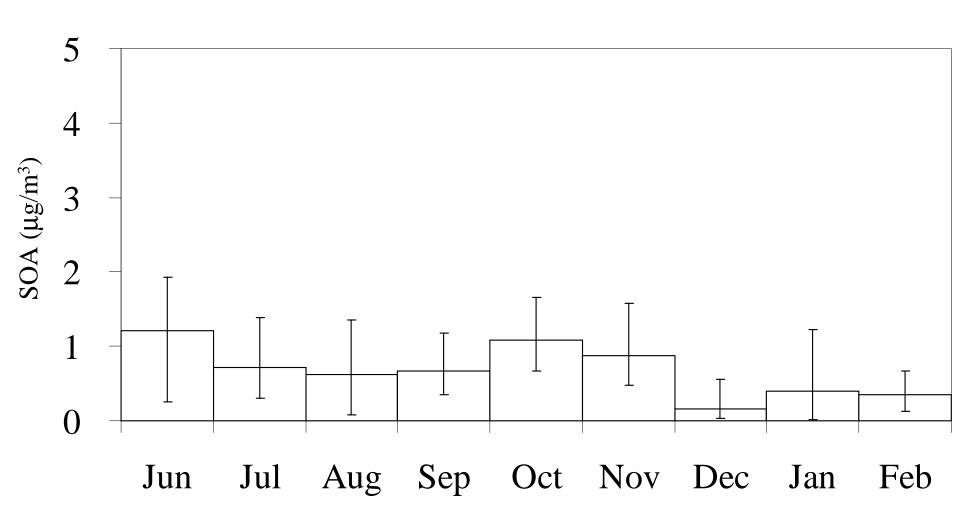
### OC and EC measurements (front Quartz) Monthly Averages (2001-2002)





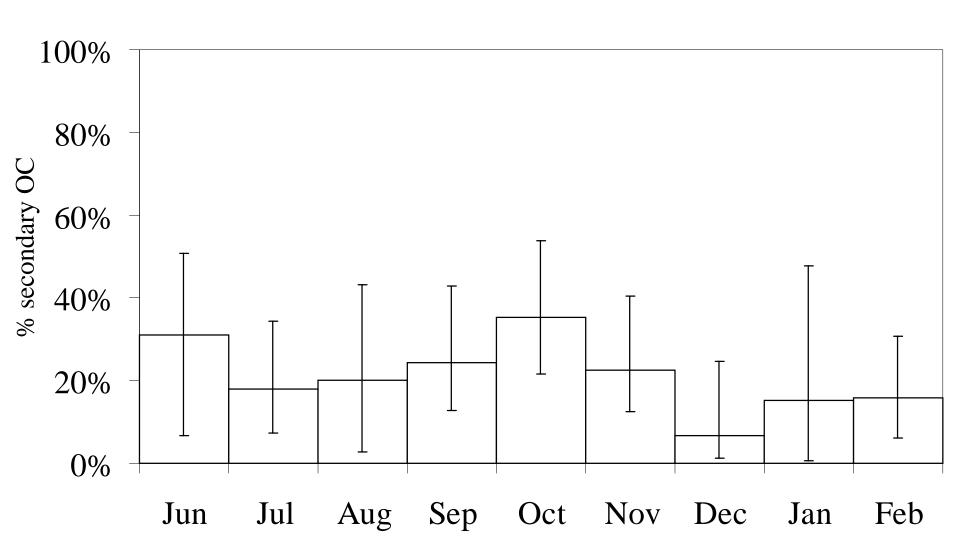


### OC and EC measurements (front Quartz) Monthly Averages (2001-2002)





## OC Composition (front Quartz), Monthly Averages (2001-2002)



### Conclusions

- More than 80% of carbonaceous material in Pittsburgh is caused by long range transport.
- 20% to 50% of OC concentration could be secondary in origin.
- Different approaches for OC/EC ratio measurement give relatively consistent results.
- Higher sampling frequency gives higher estimates of SOA. (Ability to identify periods of primary and secondary OC)





## Acknowledgements

• DOE/NETL and EPA Supersites Program for Supporting this work.